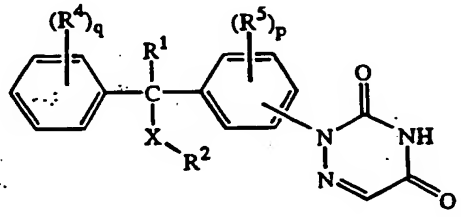
ABSTRACT

NOVEL IL-5 INHIBITING 6-AZAURACIL DERIVATIVES

The present invention is concerned with the compounds of formula



(I)

the *N*-oxides, the pharmaceutically acceptable addition salts and the stereochemically isomeric forms thereof, p and q are 0, 1, 2, 3 or 4 and q is also 5; X is O, S, NR^3 or a direct bond; R^1 is hydrogen, hydroxy, halo, optionally substituted amino, optionally substituted C_{1-6}alkyl, C_{1-6}alkyloxy, C_{3-7}cycloalkyl or aryl; R^2 is aryl, Het^1, C_{3-7}cycloalkyl, optionally substituted C_{1-6}alkyl; and if X is O, S or NR^3, then R^2 may also be a carbonyl or thiocarbonyl linked substituent; R^3 is hydrogen or C_{1-4}alkyl; R^4 and R^5 independently are optionally substituted C_{1-6}alkyl, halo, hydroxy, mercapto, C_{1-6}alkyloxy, C_{1-6}alkylthio, C_{1-6}alkylcarbonyloxy, aryl, cyano, nitro, Het^3, R^6 or NR^7R^8; R^6 is substituted sulfonyl or sulfinyl; R^7 and R^8 are hydrogen, optionally substituted C_{1-4}alkyl, aryl, a carbonyl or thiocarbonyl linked substituent, C_{3-7}cycloalkyl, Het^3 and R^6; R^9 and R^{10} are each independently selected from hydrogen, optionally substituted C_{1-4}alkyl, phenyl, a carbonyl or thiocarbonyl linked substituent, C_{3-7}cycloalkyl, Het^3 and R^6; R^{11} is hydroxy, mercapto, cyano, nitro, halo, trihalomethyl, C_{1-4}alkyloxy, carboxyl, C_{1-4}alkyloxycarbonyl, trihaloC_{1-4}alkylsulfonyloxy, R^6, NR^7R^8, $C(=O)NR^7R^8$, aryl, aryloxy, arylcarbonyl, C_{3-7}cycloalkyl, C_{3-7}cycloalkyloxy, phthalimide-2-yl, Het^3 and $C(=O)Het^3$; R^{12} and R^{13} are each independently selected from hydrogen, optionally substituted C_{1-4}alkyl, phenyl, a carbonyl or thiocarbonyl linked substituent, C_{3-7}cycloalkyl and R^6; aryl is optionally substituted phenyl; Het^1, Het^2 and Het^3 are optionally substituted heterocycles ; to processes for their preparation and compositions comprising them. It further relates to their use as a medicine.